

# Accelerating Life Sciences with Imaging Real-World Data

Investor Presentation



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## **OMN Snapshot**

## Real-World Data Industry Leader

#### NASDAQ: ONMD

- Providing innovative solutions that unlock the significant value contained in the Real-World Data ("RWD") repositories of 121.2M clinical exams from 1,413 healthcare system and provider sites via a revenue sharing model
- Incomparable in-house RWD Curation Team
  - 300+ years of clinical & radiology experience; working in all 50 states and 8+ countries
  - Certified in every imaging modality: MR, CT, IR, Cath lab, OR, Specials, US, MG, PET, NM, X-ray
- iRWD<sup>™</sup> network enables secure, comprehensive management of diverse clinical data types, including:
  - Electronic Health Records
  - Laboratory & Diagnostic Results
  - Medical Imaging Data

## Unlocking the Potential of Medical Imaging as RWD The Opportunity



FDA defines RWD as health data sourced from various electronic records, claims, registries, and digital health technologies

Medical imaging is a powerful type of RWD, but leveraging it for life sciences R&D has been challenging

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These images are large and complex objects that provide rich information; often in 3D, and often with a 4<sup>th</sup> longitudinal dimension

3

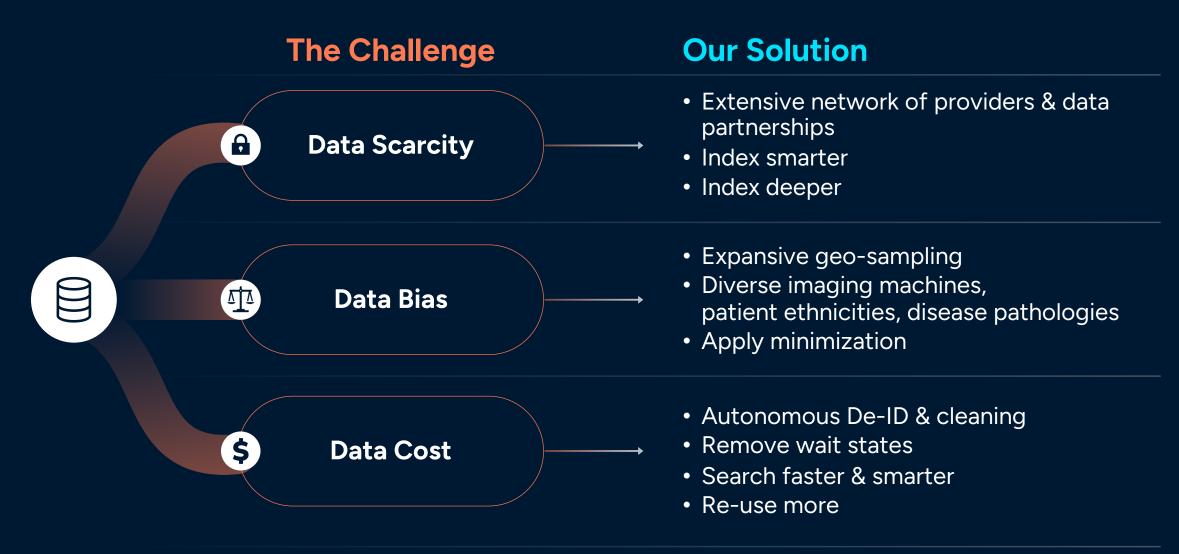
Centralizing, de-identifying, searching, and curating this RWD can yield crucial insights for life sciences organizations

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However, this requires the ability to manage petabytes of data – representing millions of patient exams and many data points

## Solving RWD's Unmet Need





## Target Markets



1	Imaging AI	Accelerate Al Model Development	<ul> <li>RWD replaces clinical trials, speeding up AI model development</li> <li>Simultaneously pursue multiple use cases with rich, heterogeneous, FDA-grade data</li> <li>Data annotation expenses are optimized based on data curation and longitudinal records that open door for weak labeling</li> </ul>
2	Drug Development	Supplement Clinical Trials with Evidence & Assess Unmet Needs	<ul> <li>RWD serves as surrogate endpoint to an outcome &amp; study control, based on phenotypical evidence</li> <li>This data must explicitly match very precise and consequential cohort specifications, and stand up the rigors of prospective clinical trials</li> <li>Willingness to pay a premium for near on-demand access vs multi-year clinical trial timelines</li> </ul>
3	Devices & Medical Implants	Product Development & Post Market Surveillance	<ul> <li>RWD used by device manufacturers to better understand the union rate of their devices</li> <li>FDA requires post market data on complication rates, removal and revision, infection rates and time to union on the device</li> <li>Willingness to pay a premium for near on-demand access for surveillance of time.</li> </ul>

## Bridging the Gap – For Analytics Needs

#### **Clinical Data**

**Medical Images** 

Electronic Medical Records (EMR)

Data from Medical Devices

Longitudinal Patient Records Reduces data delivery from **months to weeks** 

# ONEMEDNET

Only the **RIGHT DATA** at the **RIGHT TIME** in the **RIGHT FORMAT** 

#### Life Sciences Organizations

#### **Drug Development**

Existing clinical data can be used to cut costs and speed up trials

#### Medical Device Development

Post-market surveillance data is essential for FDA compliance

#### Imaging/Diagnostic Al Development

Large datasets needed to train models

## Our Solution: iRWD<sup>™</sup> Network & Platform





#### **Enables our customers to:**



Securely search, de-identify, and curate current clinical data in real-time



Create ease of access to actionable data from healthcare providers to life sciences organizations



Impact patient outcomes and streamline healthcare innovation

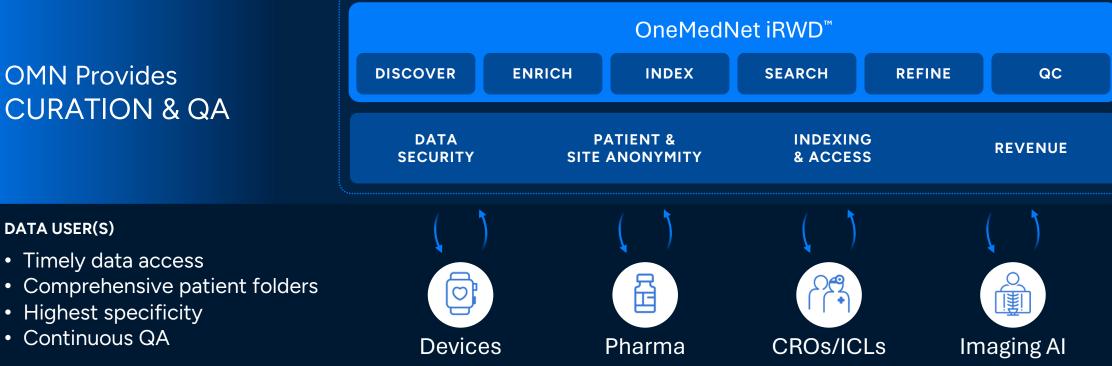
## iRWD<sup>™</sup> Federated Network



#### **OMN** Provides **CURATION & QA**

DATA USER(S)

Continuous QA



## The Difference is in the Data





#### Our Data is Regulatory-grade

- RWD is used to complement, reduce, and in some cases, replace clinical trials
- Meets strict requirements for analysis and regulatory approval
- Adds underserved populations in research
- Supporting stages from Discovery through Post-Market Surveillance



#### **Our Data Meets Complex & Precise Specifications**

- We ensures exact compliance with cohort protocols and delivery formats
- Extensive data coverage includes longitudinal patient records
- Network for ongoing imaging monitoring and patient information retrieval
- Accessing our native provider data offers comprehensive resources for initial and follow-up studies

## Superior Data Begins with a Superior Network

#### Our Ever-expanding Real-time, RWD

121M+ Clinical Exams 31M+ Patients

1,400+ Healthcare Partner Sites



Why It Matters?

#### Impact on Research:

- Diverse and comprehensive data leads to better research outcomes
- Ability to monitor and improve medical devices and treatments

## **Competitive Advantages**

demand

Modern DevOps CI/CD –

latest tools for automating

software rollout to cloud





#### **Highly Qualified Curation Staff**

- Interpreted sophisticated requirements of the buyer
- Accurately fill complex orders quickly
- Embedding learnings into the platform as AI & ML

## Proprietary, Autonomous De-Identification Process



	De-ID without Al (Others)	De-ID with AI (OneMedNet)	
Multiple De-ID Process in Parallel	Some automation but mainly limited to human curation team	Autonomous De-ID process can scale with needs. Human in the loop for Auditing	<ul> <li>Deep Learning &amp; Large Language Models (LLMs) automates the De-ID process with human in the loop for auditing</li> </ul>
De-ID Complete Study	Hours	Minutes	<ul> <li>Generative AI driven de-identification process to efficiently remove PHI in just minutes, streamlining clinical imaging curation</li> </ul>

## **Organizational Compliance**

#### Privacy / Regulatory Compliance

- HIPAA-HITECH, GDPR, PIPEDA
- FDA 21 CFR Part 11

#### Cybersecurity / Risk Management

- HITRUST Certified 3rd Party Hosting/SOC
- NIST 800-53 CSF Risk Management/Controls
- RBAC, MFA, & SSO



### Case Example <u>Client Company Requests iRWD<sup>™</sup></u> For Cancer Research



#### SAMPLE REQUIREMENTS INCLUDED:

- 1,500 screening positives; 1,500 screening negatives
- Metadata for all biopsy, cytology, or histology of specimen within 12 months of initial screening
- Considerable patient report information (e.g. tumor size, lesion location)
- Race/ethnicity population ranges from multiple sites
- Specific imaging technology vendor >70%

#### **UNMATCHED DATA CURATION:**

- Converted unstructured data into structured data
- Reconciled longitudinal and disparate patient record information
- Harmonized site dependent, and year-to-year, data formatting inconsistencies
- Achieved highly accurate patient population stipulations so no "sifting" needed by analyzing company

## 3 Pillars to Success



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#### iRWD<sup>™</sup> PLATFORM

#### Accelerate Platform Development

- Increase ease of access to repositories through modern, robust platform design
- Decrease individual and federated search and curation time through AI tools



#### PARTNERS

#### Prioritize Onboarding Partners

- Continue to expand hospital and academic medical center relationships for the **right** data repositories
- Improve revenue sharing mix over time



#### CUSTOMER SUCCESS

#### Scale & Decrease Time to Revenue

 Continue to align organization for growth investing in strategic hires to fuel growth in sales and in operations

# Investment Highlights



- A leading provider of precision curated, regulatory-grade RWD
- Differentiated iRWD<sup>™</sup> platform to bring data from healthcare providers to Life Sciences organizations
- Experienced management team
- Entering a stage of anticipated rapid growth

**4,800+** Clinical Trials in 2023<sup>2</sup> 28-34%

Increase in AI Medical Imaging Market Size Annually<sup>3</sup>

NASDAQ: ONMD

<sup>1</sup> CBInsights – Future of Clinical Trials 2021 <sup>2</sup> IQVIA Institute – Global Trends in R&D 2024 <sup>3</sup> Mordor Intelligence Market Size 2024 - 2029



Clinical Trials Market<sup>1</sup>